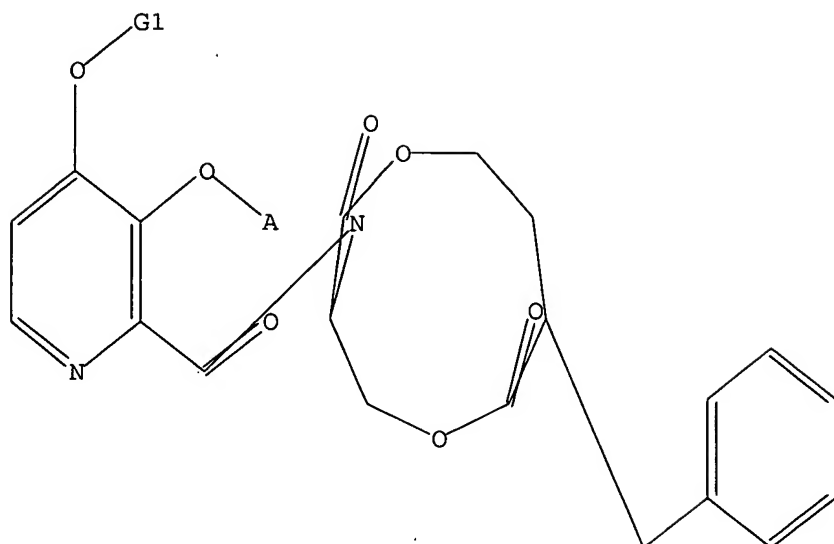


10/647,172

L17 HAS NO ANSWERS
L17 STR



G1 Me,Et,n-Pr,i-Pr,n-Bu,i-Bu,s-Bu,t-Bu

Structure attributes must be viewed using STN Express query preparation.

=> s l17 sss full
FULL SEARCH INITIATED 13:06:09 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 261 TO ITERATE

100.0% PROCESSED 261 ITERATIONS 217 ANSWERS
SEARCH TIME: 00.00.01

L18 217 SEA SSS FUL L17

=> file caplus
COST IN U.S. DOLLARS

FULL ESTIMATED COST

| SINCE FILE | TOTAL |
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| ENTRY | SESSION |
| 161.33 | 528.38 |

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

CA SUBSCRIBER PRICE

| SINCE FILE | TOTAL |
|------------|---------|
| ENTRY | SESSION |
| 0.00 | -5.11 |

FILE 'CAPLUS' ENTERED AT 13:06:22 ON 08 APR 2005
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10/647,172

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FILE COVERS 1907 - 8 Apr 2005 VOL 142 ISS 16
FILE LAST UPDATED: 7 Apr 2005 (20050407/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s l18

L19 7 L18

=> d l19 1-7 ibib abs hitstr

L19 ANSWER 1 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2003:335078 CAPLUS

DOCUMENT NUMBER: 138:337882

TITLE: Preparation of UK-2A derivatives as agricultural fungicides

INVENTOR(S): Meyer, Kevin Gerald; Rogers, Richard Brewer; Yao, Chenglin; Niyaz, Normohammed Mohamed; Adamski Butz, Jenifer Lynn; Nader, Bassam Salim

PATENT ASSIGNEE(S): Dow Agrosciences Llc Patent Department, USA

SOURCE: PCT Int. Appl., 39 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

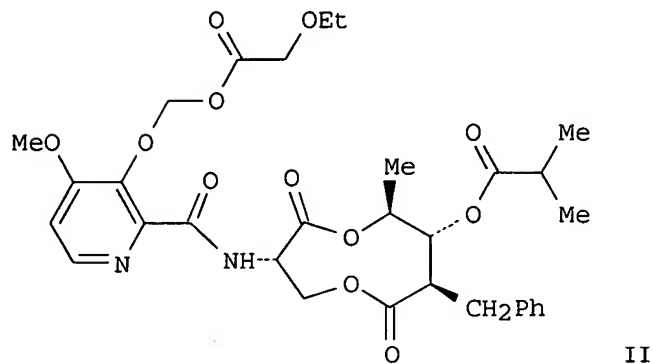
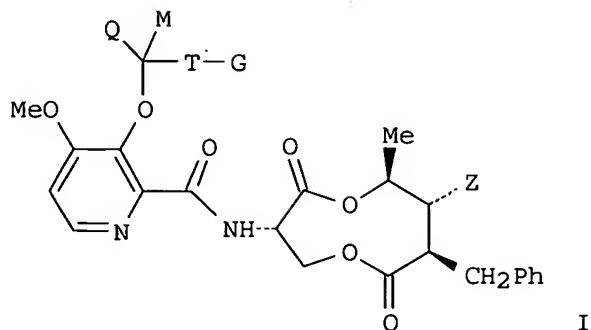
LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

Assigned

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|------------------------|--|----------|-----------------|------------|
| WO 2003035617 | A2 | 20030501 | WO 2002-US33947 | 20021023 |
| WO 2003035617 | A3 | 20031113 | | |
| W: | AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW | | | |
| RW: | GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG | | | |
| EP 1438306 | A2 | 20040721 | EP 2002-802199 | 20021023 |
| R: | AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK | | | |
| JP 2005507921 | T2 | 20050324 | JP 2003-538133 | 20021023 |
| US 2004192924 | A1 | 20040930 | US 2004-493456 | 20040423 |
| US 6861390 | B2 | 20050301 | | |
| PRIORITY APPLN. INFO.: | | | US 2001-335814P | P 20011023 |
| | | | WO 2002-US33947 | W 20021023 |
| OTHER SOURCE(S): | MARPAT 138:337882 | | | |
| GI | | | | |



AB Derivs. of UK-2A of formula I [Z = H, alkoxy, acyl, OC(O)Oalkyl, OC(O)dialkylamino, etc.; Q, M = H, Me, Et, CF₃, Ph, vinyl, cyclopropyl; T = O, OC(O), OCO₂, S, SC(O), SCO₂; G = H, alkyl, alkenyl, alkynyl, cycloalkyl, aryl, heteroaryl] are provided for the treatment of plant fungal diseases. Thus, II was prepared from UK-2A. The prepared compds. were tested for control of in vivo whole plant fungal infection.

IT 512192-31-3P 512192-33-5P 512192-36-8P
 517875-15-9P 517875-16-0P 517875-17-1P
 517875-18-2P 517875-19-3P 517875-20-6P
 517875-21-7P 517875-22-8P 517875-23-9P
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 517875-79-5P 517875-80-8P 517875-81-9P
 517875-82-0P 517875-83-1P 517875-84-2P
 517875-85-3P 517875-86-4P

RL: AGR (Agricultural use); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

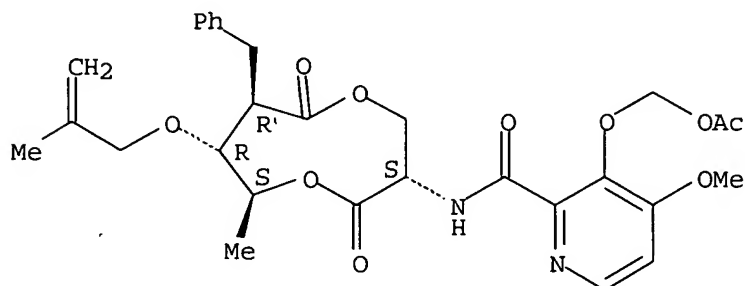
(preparation of UK-2A derivs. as agricultural fungicides)

10/647,172

RN 512192-31-3 CAPLUS

CN 2-Pyridinecarboxamide, 3-[(acetyloxy)methoxy]-4-methoxy-N-[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-2-propenyl)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]- (9CI) (CA INDEX NAME)

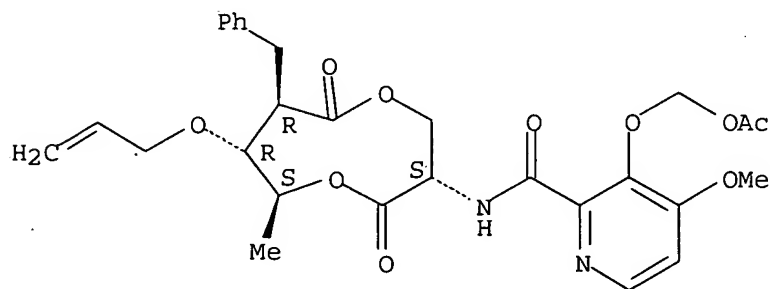
Absolute stereochemistry.



RN 512192-33-5 CAPLUS

CN 2-Pyridinecarboxamide, 3-[(acetyloxy)methoxy]-4-methoxy-N-[(3S,7R,8R,9S)-9-methyl-2,6-dioxo-7-(phenylmethyl)-8-(2-propenyloxy)-1,5-dioxonan-3-yl]- (9CI) (CA INDEX NAME)

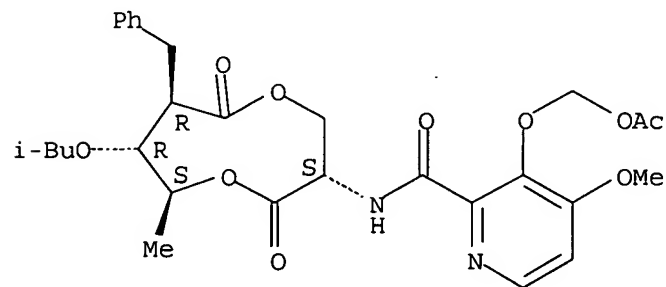
Absolute stereochemistry.



RN 512192-36-8 CAPLUS

CN 2-Pyridinecarboxamide, 3-[(acetyloxy)methoxy]-4-methoxy-N-[(3S,7R,8R,9S)-9-methyl-8-(2-methylpropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]- (9CI) (CA INDEX NAME)

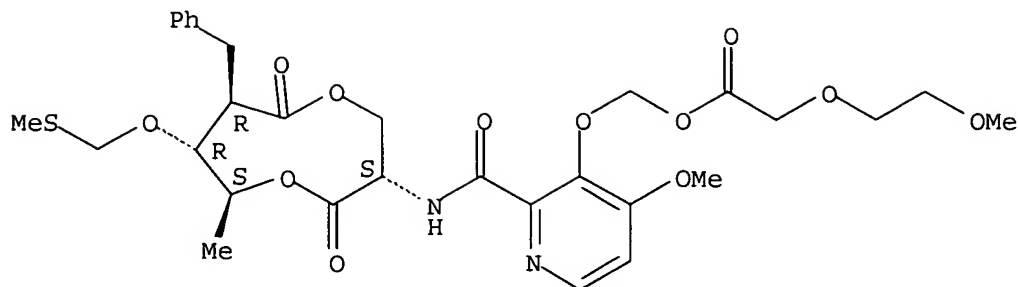
Absolute stereochemistry.



RN 517875-15-9 CAPLUS

CN Propanoic acid, 2-methyl-, (3S,6S,7R,8R)-3-[[[4-methoxy-3-(methoxymethoxy)-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

10/647,172

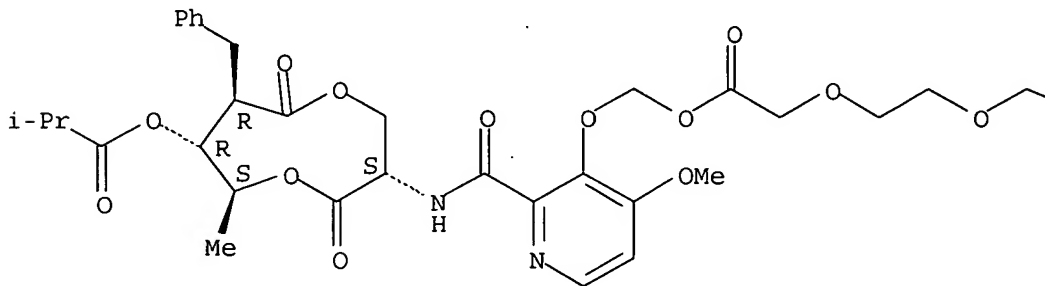


RN 517875-86-4 CAPLUS

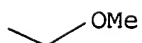
CN Propanoic acid, 2-methyl-, (3S,6S,7R,8R)-3-[[[4-methoxy-3-[(3-oxo-2,5,8,11-tetraoxadodec-1-yl)oxy]-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A



PAGE 1-B



L19 ANSWER 2 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2003:301046 CAPLUS

DOCUMENT NUMBER: 138:321054

TITLE: Process to produce alkyl-ether derivatives of UK-2A
INVENTOR(S): Niyaz, Normohammed Mohamed; Deamicis, Carl Vincent;
Rogers, Richard Brewer; Meyer, Kevin Gerald; Dent,
William Hunter, III; Anzeveno, Peter Biagio

PATENT ASSIGNEE(S): Dow Agrosciences LLC, USA

SOURCE: PCT Int. Appl., 20 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---------------|------|----------|-----------------|----------|
| WO 2003031403 | A2 | 20030417 | WO 2002-US31848 | 20021004 |
| WO 2003031403 | A3 | 20030918 | | |

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RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

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US 2004-491978

20040405

PRIORITY APPLN. INFO.:

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P 20011005

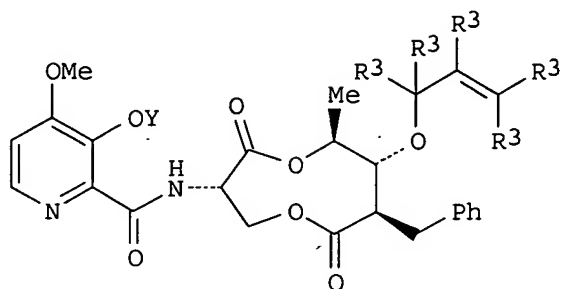
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W 20021004

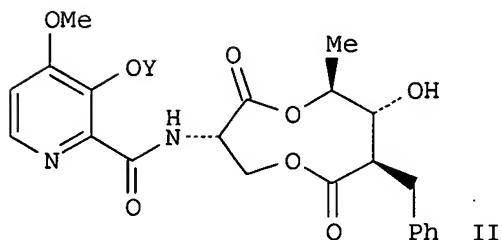
OTHER SOURCE(S):

MARPAT 138:321054

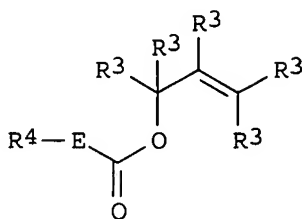
GI



I



II



III

AB A process is disclosed for the preparation of allyl-alkyl ether derivs. I [Y = H, benzyl, Si(alkyl)₃, etc.; R₃ = H, alk(en/yn)yl, cycloalkyl, (hetero)aryl] of antibiotic UK-2A. The process is comprised of coupling II with III [E = O, NR₆; R₄, R₆ = alkyl, aryl] in the presence of a catalyst complex and solvent. For instance II [Y = PhCH₂] was coupled to Et methallylcarbonate (dppf, Pd₂dba₃) to give the corresponding methallyl derivative of I. Several examples are provided and subsequent sidechain reduction is also described.

IT 496781-72-7P 512192-28-8P 512192-29-9P

512192-30-2P 512192-31-3P 512192-32-4P

512192-33-5P 512192-34-6P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

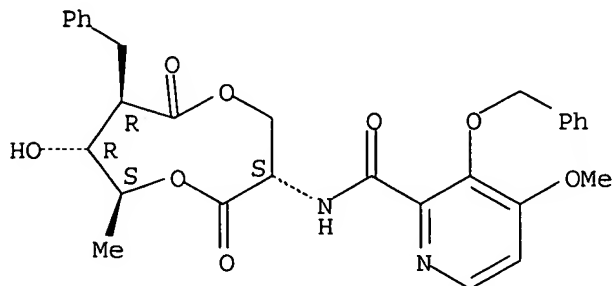
(palladium catalyzed allylation process to produce alkyl-ether derivs. of UK-2A)

RN 496781-72-7 CAPLUS

10/647,172

CN 2-Pyridinecarboxamide, N-[(3S,7R,8R,9S)-8-hydroxy-9-methyl-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]-4-methoxy-3-(phenylmethoxy)- (9CI) (CA INDEX NAME)

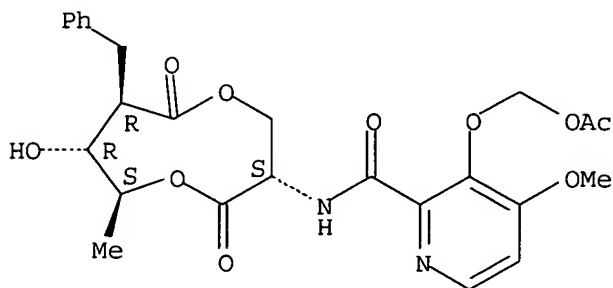
Absolute stereochemistry.



RN 512192-28-8 CAPLUS

CN 2-Pyridinecarboxamide, 3-[(acetyloxy)methoxy]-N-[(3S,7R,8R,9S)-8-hydroxy-9-methyl-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]-4-methoxy- (9CI) (CA INDEX NAME)

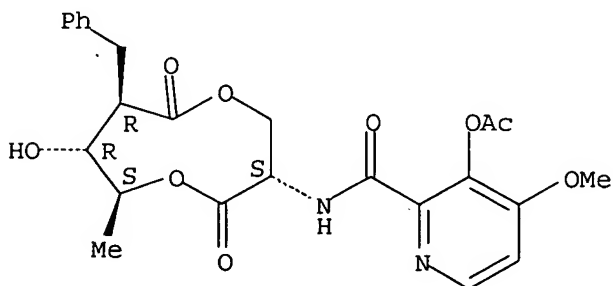
Absolute stereochemistry.



RN 512192-29-9 CAPLUS

CN 2-Pyridinecarboxamide, 3-(acetyloxy)-N-[(3S,7R,8R,9S)-8-hydroxy-9-methyl-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]-4-methoxy- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

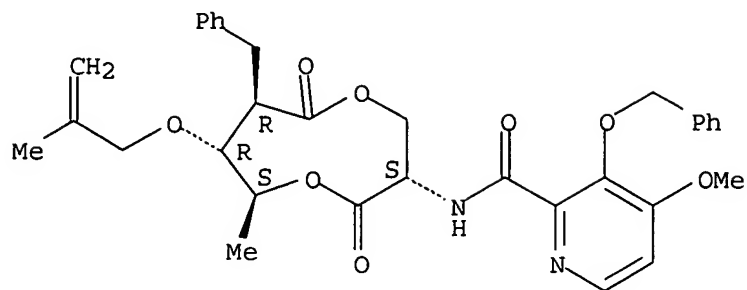


RN 512192-30-2 CAPLUS

CN 2-Pyridinecarboxamide, 4-methoxy-N-[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-2-propenyl)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]-3-(phenylmethoxy)- (9CI) (CA INDEX NAME)

10/647,172

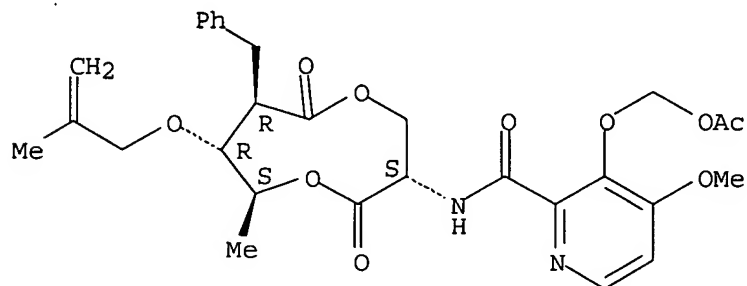
Absolute stereochemistry.



RN 512192-31-3 CAPLUS

CN 2-Pyridinecarboxamide, 3-[(acetyloxy)methoxy]-4-methoxy-N-[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-2-propenyl)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]- (9CI) (CA INDEX NAME)

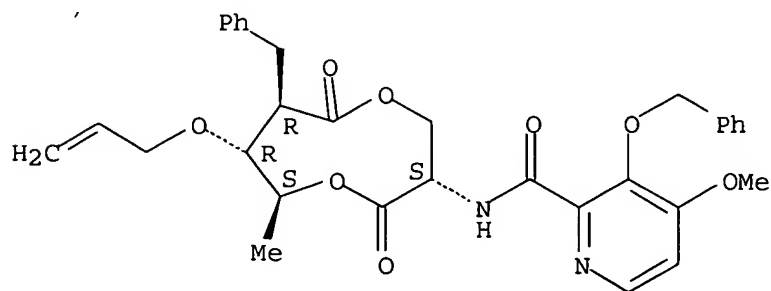
Absolute stereochemistry.



RN 512192-32-4 CAPLUS

CN 2-Pyridinecarboxamide, 4-methoxy-N-[(3S,7R,8R,9S)-9-methyl-2,6-dioxo-7-(phenylmethyl)-8-(2-propenyloxy)-1,5-dioxonan-3-yl]-3-(phenylmethoxy)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

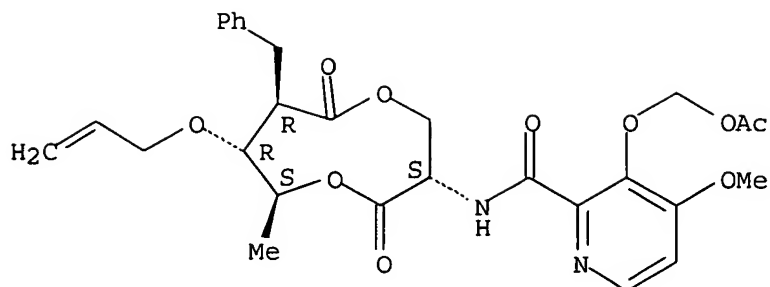


RN 512192-33-5 CAPLUS

CN 2-Pyridinecarboxamide, 3-[(acetyloxy)methoxy]-4-methoxy-N-[(3S,7R,8R,9S)-9-methyl-2,6-dioxo-7-(phenylmethyl)-8-(2-propenyloxy)-1,5-dioxonan-3-yl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

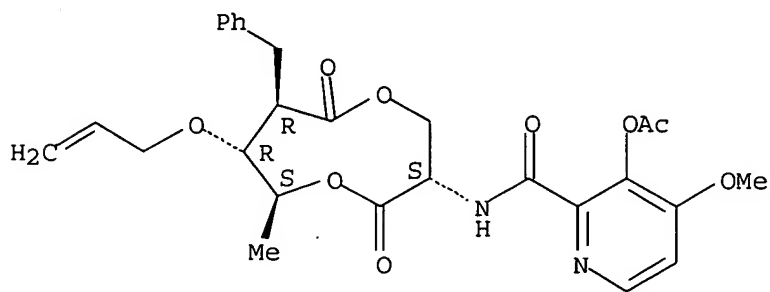
10/647,172



RN 512192-34-6 CAPLUS

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(CA INDEX NAME)

Absolute stereochemistry.



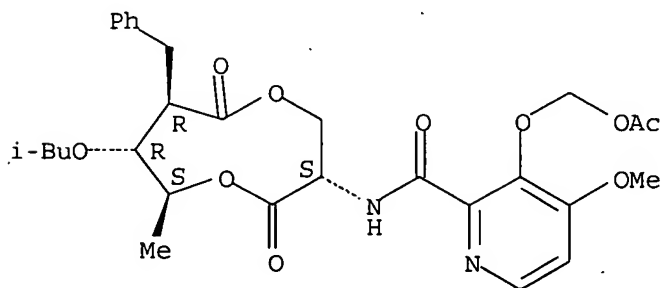
IT 512192-36-8P 512192-38-0P

RL: SPN (Synthetic preparation); PREP (Preparation)
(palladium catalyzed allylation process to produce alkyl-ether derivs.
of UK-2A)

RN 512192-36-8 CAPLUS

CN 2-Pyridinecarboxamide, 3-[(acetyloxy)methoxy]-4-methoxy-N-[(3S,7R,8R,9S)-9-methyl-8-(2-methylpropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

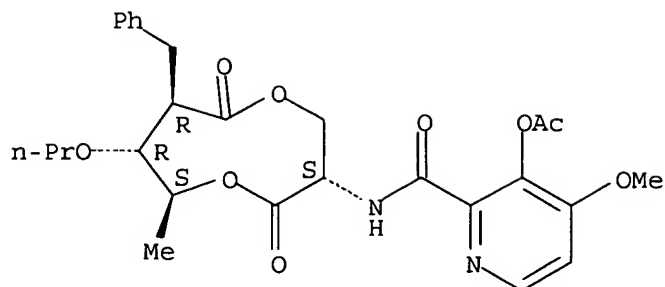


RN 512192-38-0 CAPLUS

CN 2-Pyridinecarboxamide, 3-(acetyloxy)-4-methoxy-N-[(3S,7R,8R,9S)-9-methyl-2,6-dioxo-7-(phenylmethyl)-8-propoxy-1,5-dioxonan-3-yl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

10/647,172



L19 ANSWER 3 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2003:117821 CAPLUS

DOCUMENT NUMBER: 138:153370

TITLE: Preparation of UK-2A derivatives via reductive cleavage of the exocyclic ester of UK-2A or its derivatives

INVENTOR(S): Meyer, Kevin Gerald; Niyaz, Normohammed Mohamed; Deamicis, Carl Vincent; Rogers, Richard Brewer

PATENT ASSIGNEE(S): Dow Agrosciences LLC, USA

SOURCE: PCT Int. Appl., 15 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

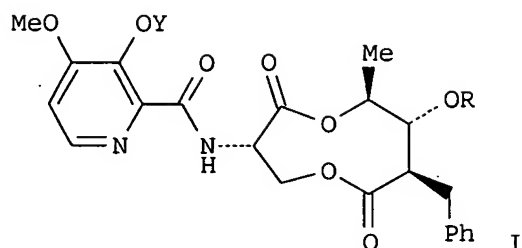
LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

Assigned

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|------------------------|--|--|-----------------|------------|
| WO 2003011857 | A1 | 20030213 | WO 2002-US24204 | 20020731 |
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| BR 2002011534 | A | 20040713 | BR 2002-11534 | 20020731 |
| JP 2005501836 | T2 | 20050120 | JP 2003-517049 | 20020731 |
| US 2004171838 | A1 | 20040902 | US 2004-483947 | 20040115 |
| PRIORITY APPLN. INFO.: | | | US 2001-308939P | P 20010731 |
| | | | WO 2002-US24204 | W 20020731 |
| OTHER SOURCE(S): | | CASREACT 138:153370; MARPAT 138:153370 | | |
| GI | | | | |



AB The present invention discloses a process for the preparation of UK-2A derivs., such as I [R = H; Y = H, (un)substituted benzyl, CH₂OC₁₋₈ alkyl, CH₂OC₃₋₈ cycloalkyl, allyl, (un)substituted tetrahydropyranyl, (un)substituted tetrahydrofuranyl, Si(C₁₋₄ alkyl)₃, and Si(Ph)_x(C₁₋₄ alkyl)_{3-x} where x = 1-3], via reductive cleavage of the exocyclic ester of UK-2A I [R = OCOCH(Me)₂; Y = H (II)] or its derivs., such as I [R = COCH(Me)₂; Y = H, (un)substituted benzyl, CH₂OC₁₋₈ alkyl, CH₂OC₃₋₈ cycloalkyl, allyl, (un)substituted tetrahydropyranyl, (un)substituted tetrahydrofuranyl, Si(C₁₋₄ alkyl)₃, and Si(Ph)_x(C₁₋₄ alkyl)_{3-x} where x = 1-3], in the presence of a reducing agent and in the presence of an aprotic solvent. Thus, II was reacted with benzyl bromide to afford O-benzylated derivative I [R = OCOCH(Me)₂; Y = CH₂Ph], which was treated with diisobutylaluminum hydride to afford UK-2A derivative I [R = H; Y = CH₂Ph].

IT 234112-89-1P

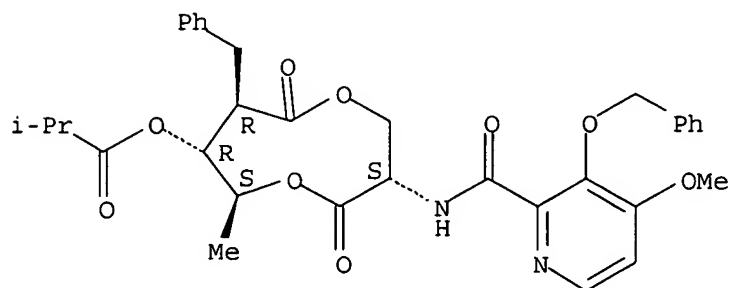
RL: IMF (Industrial manufacture); RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of UK-2A derivs. via reductive cleavage of the exocyclic ester of UK-2A or its derivs.)

RN 234112-89-1 CAPLUS

CN Propanoic acid, 2-methyl-, (3S,6S,7R,8R)-3-[[[4-methoxy-3-(phenylmethoxy)-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.



IT 496781-72-7P

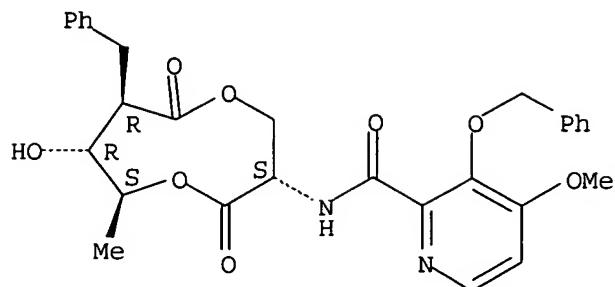
RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP (Preparation)

(preparation of UK-2A derivs. via reductive cleavage of the exocyclic ester of UK-2A or its derivs.)

RN 496781-72-7 CAPLUS

CN 2-Pyridinecarboxamide, N-[(3S,7R,8R,9S)-8-hydroxy-9-methyl-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]-4-methoxy-3-(phenylmethoxy)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L19 ANSWER 4 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2001:152650 CAPLUS

DOCUMENT NUMBER: 134:207831

TITLE: Preparation, composition and use of heterocyclic aromatic amides as fungicides

INVENTOR(S): Ricks, Michael John; Dent, William Hunter, III; Rogers, Richard Brewer; Yao, Chenglin; Nader, Bassam Salim; Miesel, John Louis; Fitzpatrick, Gina Marie; Meyer, Kevin Gerald; Niyaz, Noormohamed Mohamed; Morrison, Irene Mae; Henry, Matthew James; Adamski, Butz Jenifer Lynn; Gajewski, Robert Peter

PATENT ASSIGNEE(S): Dow Agrosciences LLC, USA

SOURCE: PCT Int. Appl., 200 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

= present

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|------|----------|-----------------|----------|
| WO 2001014339 | A2 | 20010301 | WO 2000-US21523 | 20000804 |
| WO 2001014339 | A3 | 20011115 | | |
| W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM | | | | |
| RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG | | | | |
| US 6521622 | B1 | 20030218 | US 2000-620662 | 20000720 |
| CA 2376275 | AA | 20010301 | CA 2000-2376275 | 20000804 |
| AU 2000065267 | A5 | 20010319 | AU 2000-65267 | 20000804 |
| AU 778108 | B2 | 20041118 | | |
| US 6355660 | B1 | 20020312 | US 2000-632930 | 20000804 |
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| EP 1234825 | A3 | 20030618 | | |
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| EP 1234826 | A3 | 20030618 | | |
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| EP 1234827 | A3 | 20030618 | | |
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| TR 200200409 | T2 | 20030321 | TR 2002-200200409 | 20000804 |
| BR 2000013469 | A | 20030429 | BR 2000-13469 | 20000804 |
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| EP 1486489 | A2 | 20041215 | EP 2004-22082 | 20000804 |
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| IE, FI, CY | | | | |
| EP 1493733 | A2 | 20050105 | EP 2004-22081 | 20000804 |
| EP 1493733 | A3 | 20050126 | | |
| R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, | | | | |
| IE, FI, CY | | | | |
| US 2002177578 | A1 | 20021128 | US 2001-22413 | 20011213 |
| US 2003018052 | A1 | 20030123 | US 2001-22207 | 20011213 |
| US 2003018012 | A1 | 20030123 | US 2001-22511 | 20011213 |
| US 6706740 | B2 | 20040316 | | |
| US 2003022902 | A1 | 20030130 | US 2001-22483 | 20011213 |
| US 2003022903 | A1 | 20030130 | US 2001-23497 | 20011213 |
| ZA 2002000435 | A | 20030117 | ZA 2002-435 | 20020117 |
| US 2004034025 | A1 | 20040219 | US 2002-307844 | 20021202 |
| US 2004048864 | A1 | 20040311 | US 2002-307710 | 20021202 |
| PRIORITY APPLN. INFO.: | | | US 1999-149977P | P 19990820 |
| | | | US 1999-150248P | P 19990823 |
| | | | US 2000-620662 | A 20000720 |
| | | | US 1999-144676P | P 19990720 |
| | | | EP 2000-952599 | A3 20000804 |
| | | | US 2000-632930 | A3 20000804 |
| | | | WO 2000-US21523 | W 20000804 |
| OTHER SOURCE(S): | | MARPAT 134:207831 | | |
| GI | | | | |

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB Title compds. [I; wherein X1-X4 independently = O, S, NR1, N, CR2, bond; R1 = H, C1-3 alkyl, C2-3 alkenyl, C2-3 alkynyl, OH, CHF2, C1-4 alkoxy; R2 = H, F, Cl, Br, CN, OH, C1-3 alkyl, C1-3 haloalkyl cyclopropyl, C1-3 alkoxy; Z = O, S, NOH, NOR3; R3 = C1-3 alkyl; A = C1-14 alkyl, C1-14 alkynyl, C1-14 cycloalkyl, aryl, heteroaryl, Q; M = H, Si(t-Bu)Me2, Si(Ph)Me2, SiEt3, CZR4, SO2R5; R4 = H, C1-6 alkyl, C2-6 alkenyl, C2-6 alkynyl; R5 = aryl, heteroaryl, C1-6 alkyl, C2-6 alkenyl, C3-6 alkenyl, C3-6 alkynyl, C3-6 cycloalkyl; X, Y independently = O, S; W = O, CH2, bond; R = C1-8 alkyl, C2-8 alkenyl, C2-8 alkynyl, C3-8 cycloalkyl, aryl, heteroaryl; R11 = H, C1-3 alkyl, C2-5 alkenyl, C2-5 alkynyl; R10 = H, R, OR, OCOR, OCOOR; R8, R9 independently = H, C1-6 alkyl, C2-6 alkenyl; R6, R7 independently = H, C1-6 alkyl, C2-6 alkenyl, C2-5 alkynyl, C3-6 cycloalkyl] are prepared as fungicides involving application methods of effective usage of title compds. to control fungi, particularly plant pathogens and wood decaying

fungi. The invention also encompasses hydrates, salts and complexes thereof. The title compound II was prepared and tested as fungicide.

IT 321601-47-2

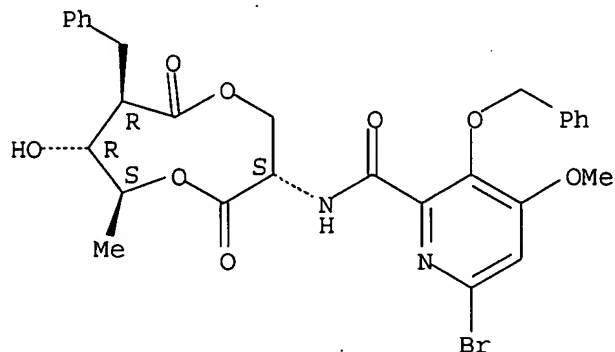
RL: RCT (Reactant); RACT (Reactant or reagent)

(preparation and fungicidal activity of heterocyclic aromatic amides)

RN 321601-47-2 CAPLUS

CN 2-Pyridinecarboxamide, 6-bromo-N-[(3S,7R,8R,9S)-8-hydroxy-9-methyl-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]-4-methoxy-3-(phenylmethoxy)-(9CI) (CA INDEX NAME)

Absolute stereochemistry.



IT 321597-69-7P 321597-70-0P 321597-71-1P

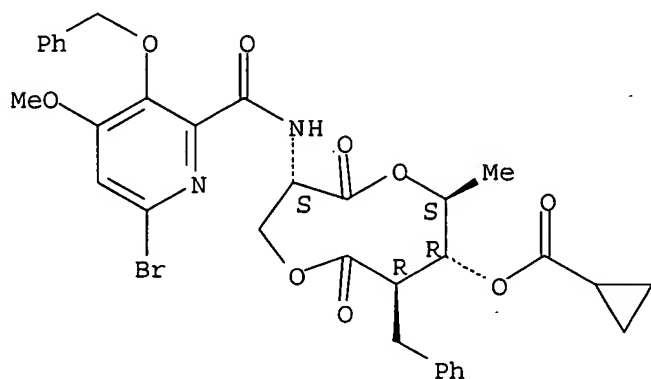
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation and fungicidal activity of heterocyclic aromatic amides)

RN 321597-69-7 CAPLUS

CN Cyclopropanecarboxylic acid, (3S,6S,7R,8R)-3-[[[6-bromo-4-methoxy-3-(phenylmethoxy)-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

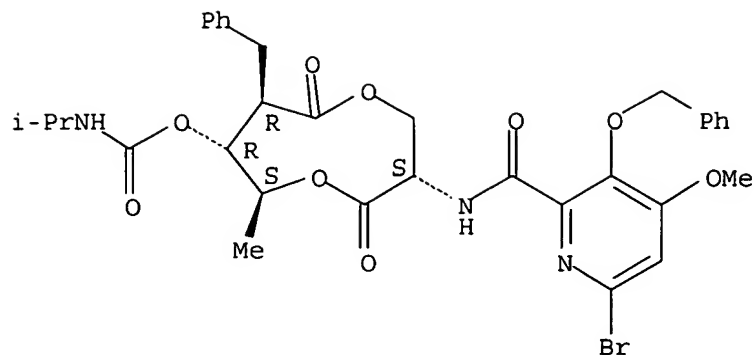


RN 321597-70-0 CAPLUS

CN Carbamic acid, (1-methylethyl)-, (3S,6S,7R,8R)-3-[[[6-bromo-4-methoxy-3-(phenylmethoxy)-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

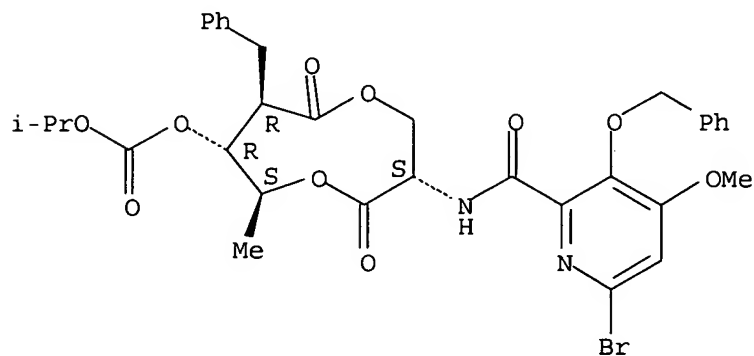
10/647,172



RN 321597-71-1 CAPLUS

CN Carbonic acid, (3S,6S,7R,8R)-3-[[[6-bromo-4-methoxy-3-(phenylmethoxy)-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl 1-methylethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.



IT 328255-87-4P 328255-88-5P 328255-89-6P
328255-90-9P 328255-91-0P 328255-92-1P
328255-93-2P 328255-94-3P 328255-95-4P
328255-96-5P 328255-97-6P 328256-00-4P
328256-01-5P 328256-02-6P 328256-03-7P
328256-15-1P 328256-16-2P 328256-17-3P
328256-21-9P 328256-23-1P 328256-24-2P
328256-25-3P 328256-26-4P 328256-27-5P
328256-28-6P 328256-29-7P 328256-31-1P
328256-32-2P 328256-33-3P 328256-36-6P
328256-37-7P 328256-38-8P 328256-39-9P
328256-40-2P 328256-42-4P 328256-45-7P
328256-47-9P 328256-56-0P 328256-57-1P
328256-58-2P 328256-59-3P 328256-60-6P
328256-61-7P 328256-62-8P 328256-63-9P
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328256-67-3P 328256-68-4P 328256-76-4P
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328256-85-5P 328256-86-6P 328256-87-7P
328256-88-8P 328256-89-9P 328256-91-3P
328257-06-3P 328257-07-4P 328257-08-5P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)

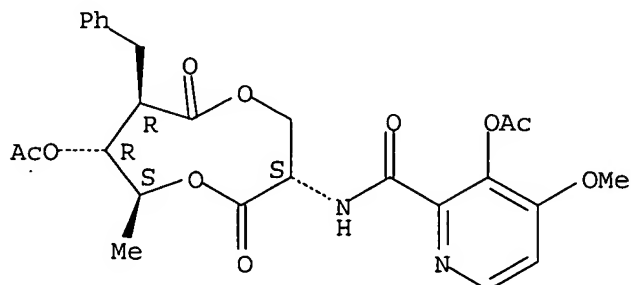
(preparation of heterocyclic aromatic amides as fungicides)

RN 328255-87-4 CAPLUS

10/647,172

CN 2-Pyridinecarboxamide, 3-(acetyloxy)-N-[(3S,7R,8R,9S)-8-(acetyloxy)-9-methyl-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]-4-methoxy- (9CI) (CA INDEX NAME)

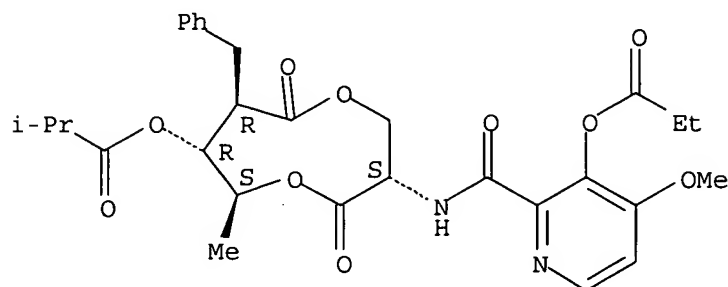
Absolute stereochemistry.



RN 328255-88-5 CAPLUS

CN Propanoic acid, 2-methyl-, (3S,6S,7R,8R)-3-[[[4-methoxy-3-(1-oxopropoxy)-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

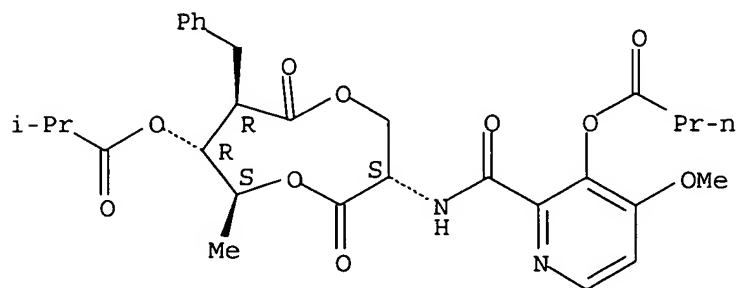
Absolute stereochemistry.



RN 328255-89-6 CAPLUS

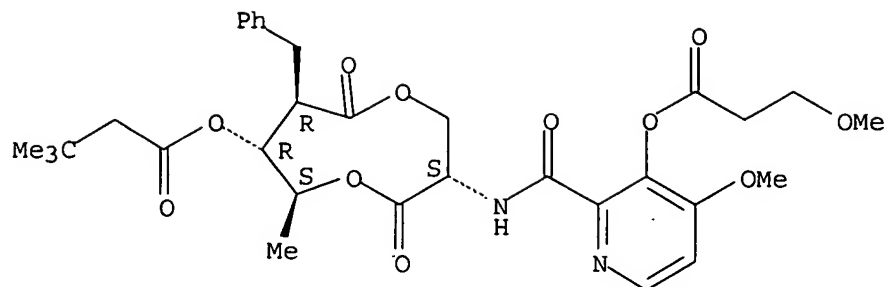
CN Butanoic acid, 4-methoxy-2-[[[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-1-oxopropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]amino]carbonyl]-3-pyridinyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 328255-90-9 CAPLUS

CN 2-Butenoic acid, 4-methoxy-2-[[[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-1-oxopropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]amino]carbonyl]-3-pyridinyl ester, (2E)- (9CI) (CA INDEX NAME)



L19 ANSWER 5 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2001:63978 CAPLUS

DOCUMENT NUMBER: 134:131431

TITLE: Fungicidal heterocyclic aromatic amides and their compositions, methods of use and preparation

INVENTOR(S): Ricks, Michael John; Dent, William Hunter, III; Rogers, Richard Brewer; Yao, Chenglin; Nader, Bassam Salim; Miesel, John Louis; Fitzpatrick, Gina Marie; Meyer, Kevin Gerald; Niyaz, Noormohamed Mohamed; Morrison, Irene Mae; Gajewski, Robert Peter

PATENT ASSIGNEE(S): Dow Agrosciences LLC, USA

SOURCE: PCT Int. Appl., 159 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

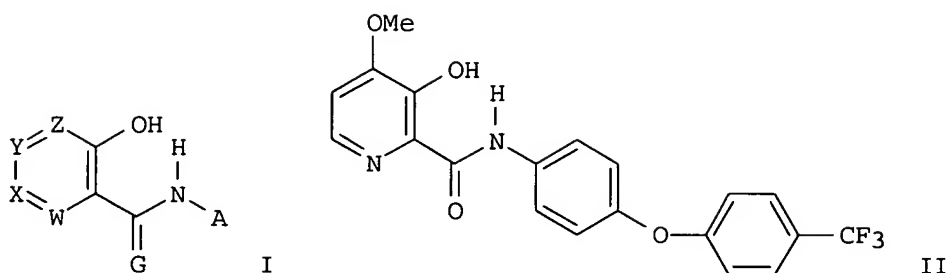
PATENT INFORMATION:

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| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---------------|--|----------|-------------------|----------|
| WO 2001005769 | A2 | 20010125 | WO 2000-US19794 | 20000720 |
| WO 2001005769 | A3 | 20011122 | | |
| W: | AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM | | | |
| RW: | GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG | | | |
| CA 2374995 | AA | 20010125 | CA 2000-2374995 | 20000720 |
| EP 1196388 | A2 | 20020417 | EP 2000-950470 | 20000720 |
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| JP 2003528806 | T2 | 20030930 | JP 2001-511430 | 20000720 |
| BR 2000012615 | A | 20040330 | BR 2000-12615 | 20000720 |
| TR 200200587 | T2 | 20041221 | TR 2002-200200587 | 20000720 |
| EP 1516874 | A1 | 20050323 | EP 2004-27006 | 20000720 |
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| EP 1516875 | A1 | 20050323 | EP 2004-27015 | 20000720 |
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| US 6355660 | B1 | 20020312 | US 2000-632930 | 20000804 |
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| US 2003018052 | A1 | 20030123 | US 2001-22207 | 20011213 |
| US 2003018012 | A1 | 20030123 | US 2001-22511 | 20011213 |

| | | | | |
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| US 6706740 | B2 | 20040316 | | |
| US 2003022902 | A1 | 20030130 | US 2001-22483 | 20011213 |
| US 2003022903 | A1 | 20030130 | US 2001-23497 | 20011213 |
| ZA 2002000436 | A | 20040302 | ZA 2002-436 | 20020117 |
| US 2004034025 | A1 | 20040219 | US 2002-307844 | 20021202 |
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| PRIORITY APPLN. INFO.: | | | US 1999-144676P | P 19990720 |
| | | | US 1999-149977P | P 19990820 |
| | | | US 1999-150248P | P 19990823 |
| | | | EP 2000-950470 | A3 20000720 |
| | | | US 2000-620662 | A3 20000720 |
| | | | WO 2000-US19794 | W 20000720 |
| | | | US 2000-632930 | A3 20000804 |

OTHER SOURCE(S): MARPAT 134:131431
GI



AB Title compds. I [W, X, Y, Z are selected from S, O, NR1, N, CR2 or bond and comprise a 5-6 membered (un)substituted heterocyclic ring; R1 = H, alkyl, alkenyl, alkynyl, OH, acyloxy, alkoxy, methyl, CHF2, cyclopropyl, or alkoxy; R2 = H, halo, CN, OH, alkyl, haloalkyl, cyclopropyl, alkoxy, haloalkoxy, etc.; G = O, S or NOR3 where R3 = H or alkyl; A = (un)substituted alkyl, alkenyl, alkynyl, cycloalkyl, unsatd. cycloalkyl, heterocycle, bi or tricyclic ring system which may contain heteroatoms, aryl or heteroaryl, etc.] bearing a hydroxy group adjacent to the amide functionality are prepared and disclosed as antifungal agents, particularly for plants. Thus, pyridinyl carboxamide II was prepared via amidation of 3-benzyloxy-6-bromo-4-methoxypyridin-2-carbonyl chloride with 4-(4-trifluoromethylphenoxy)aniline with subsequent deprotection. The preferred fungicidal composition consists of a compound of formula I with a phytol. acceptable carrier. Activity has been demonstrated against a variety of fungi, e.g., *Plasmopara viticola* (Downy Mildew of Grape), *Phytophthora infestans* (Late Blight of Tomato), and *Venturia inaequalis* (Apple Scab). I is both useful for eradication and prevention of fungal attack.

IT 321601-47-2

RL: RCT (Reactant); RACT (Reactant or reagent)

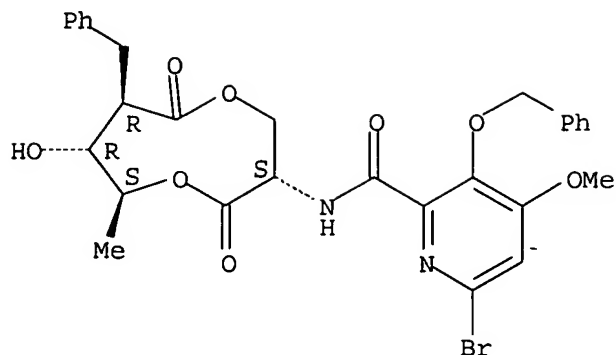
(preparation and fungicidal activity of heterocyclic aromatic amides)

RN 321601-47-2 CAPLUS

CN 2-Pyridinecarboxamide, 6-bromo-N-[(3S,7R,8R,9S)-8-hydroxy-9-methyl-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]-4-methoxy-3-(phenylmethoxy)-(9CI) (CA INDEX NAME)

Absolute stereochemistry.

10/647,172



IT 321597-69-7P 321597-70-0P 321597-71-1P

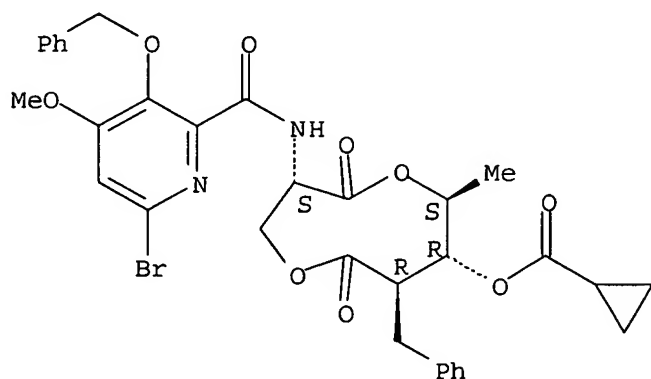
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation and fungicidal activity of heterocyclic aromatic amides)

RN 321597-69-7 CAPLUS

CN Cyclopropanecarboxylic acid, (3S,6S,7R,8R)-3-[[[6-bromo-4-methoxy-3-(phenylmethoxy)-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

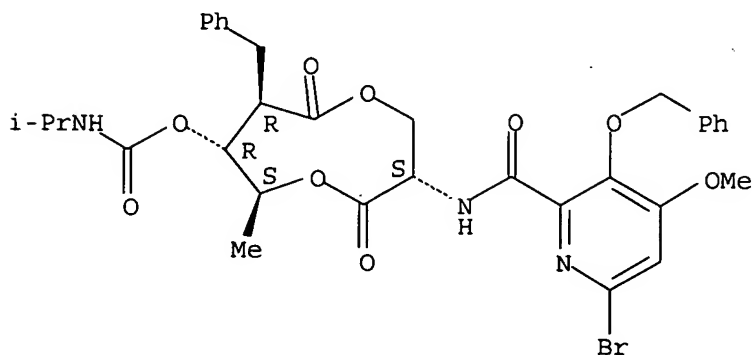
Absolute stereochemistry.



RN 321597-70-0 CAPLUS

CN Carbamic acid, (1-methylethyl)-, (3S,6S,7R,8R)-3-[[[6-bromo-4-methoxy-3-(phenylmethoxy)-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

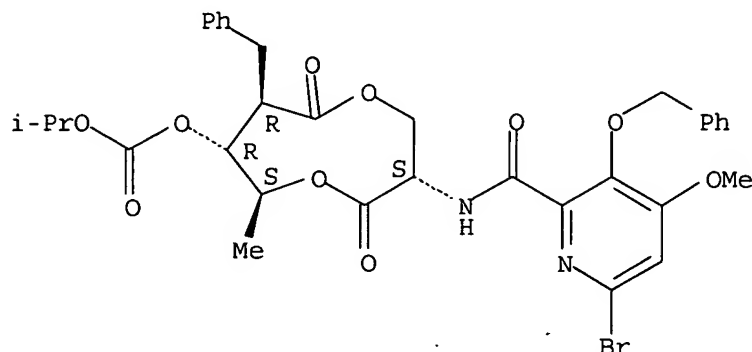


10/647,172

RN 321597-71-1 CAPLUS

CN Carbonic acid, (3S,6S,7R,8R)-3-[[[6-bromo-4-methoxy-3-(phenylmethoxy)-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl 1-methylethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L19 ANSWER 6 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 1999:511149 CAPLUS

DOCUMENT NUMBER: 131:129825

TITLE: Novel antifungal compounds and process for producing the same

INVENTOR(S): Sakanaka, Osamu; Teraoka, Takeshi; Mitomo, Koichi; Tamura, Takayoshi; Murai, Yasushi; Iinuma, Katsuharu; Kuzuhara, Kikuko; Mikoshiba, Haruki; Taniguchi, Makoto

PATENT ASSIGNEE(S): Meiji Seika Kaisha, Ltd., Japan

SOURCE: PCT Int. Appl., 92 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

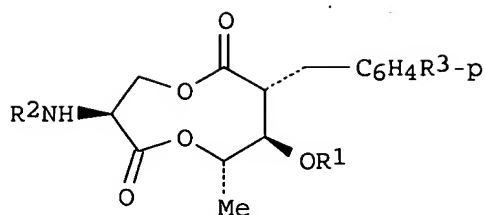
LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

data not good

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|------------------------|--|----------|-----------------|------------|
| WO 9940081 | A1 | 19990812 | WO 1999-JP541 | 19990208 |
| W: | AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM | | | |
| RW: | GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG | | | |
| CA 2319807 | AA | 19990812 | CA 1999-2319807 | 19990208 |
| AU 9924398 | A1 | 19990823 | AU 1999-24398 | 19990208 |
| AU 751098 | B2 | 20020808 | | |
| EP 1054011 | A1 | 20001122 | EP 1999-903901 | 19990208 |
| R: | AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI | | | |
| NZ 506249 | A | 20030429 | NZ 1999-506249 | 19990208 |
| PRIORITY APPLN. INFO.: | | | JP 1998-26257 | A 19980206 |
| | | | WO 1999-JP541 | W 19990208 |
| OTHER SOURCE(S): | MARPAT 131:129825 | | | |
| GI | | | | |



I

AB The title compds. [I; R1 = iso-Bu, tigloyl, isovaleryl, 2-methylbutanoyl; R2 = H, aromatic acyl, protecting group such substituted benzoyl, substituted nicotinoyl; R3 = H, nitro, amino, acylamino, N,N-dialkylamino; with provisos] are prepared Thus, UK-2A in CH₂Cl₂ containing pyridine and PC15 was refluxed for 1.5 h, the reaction mixture was allowed to cool and then reacted with methanol for 15 h to give (2R,3R,4S,7S)-7-amino-2-benzyl-5,9-dioxa-3-isobutyryloxy-4-methyl-1,6-cyclononanedione. In an antifungal test, (2R,3R,4S,7S)-7-(2-hydroxynicotinylamino)-2-benzyl-5,9-dioxa-3-isobutyryl-4-methyl-1,6-cyclononanedione (also prepared) at 0.05 µg showed potency almost double that of UK-2A against *Saccharomyces cerevisiae*.

IT 234112-85-7P 234112-86-8P 234112-89-1P

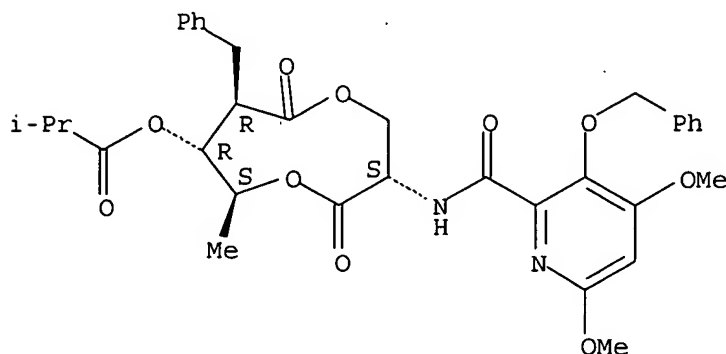
234112-90-4P 234113-05-4P 234113-06-5P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); IMF (Industrial manufacture); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)
(preparation of UK-2A derivs. as antifungals)

RN 234112-85-7 CAPLUS

CN Propanoic acid, 2-methyl-, (3S,6S,7R,8R)-3-[[[4,6-dimethoxy-3-(phenylmethoxy)-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

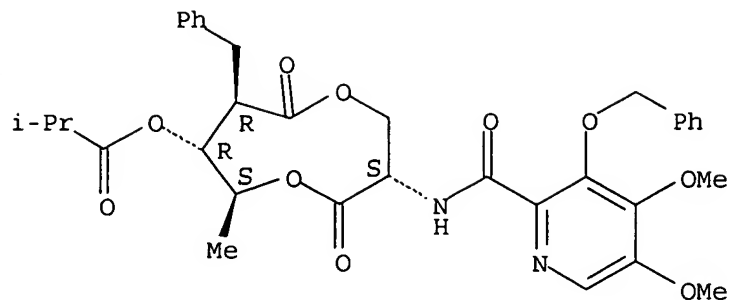


RN 234112-86-8 CAPLUS

CN Propanoic acid, 2-methyl-, (3S,6S,7R,8R)-3-[[[4,5-dimethoxy-3-(phenylmethoxy)-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

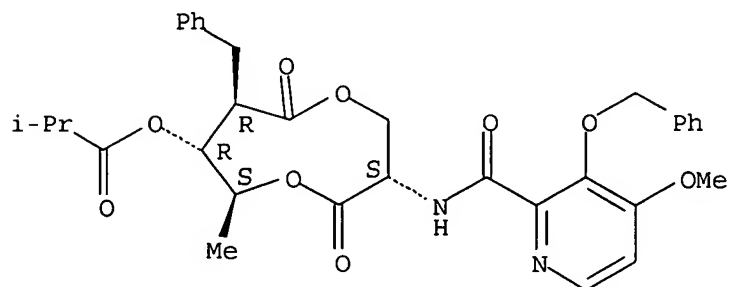
10/647,172



RN 234112-89-1 CAPLUS

CN Propanoic acid, 2-methyl-, (3S,6S,7R,8R)-3-[[[4-methoxy-3-(phenylmethoxy)-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

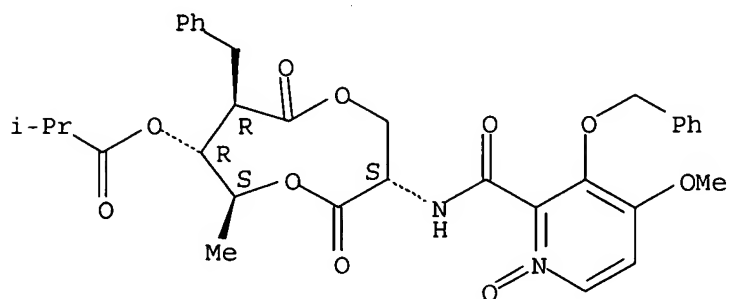
Absolute stereochemistry.



RN 234112-90-4 CAPLUS

CN Propanoic acid, 2-methyl-, (3S,6S,7R,8R)-3-[[[4-methoxy-1-oxido-3-(phenylmethoxy)-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

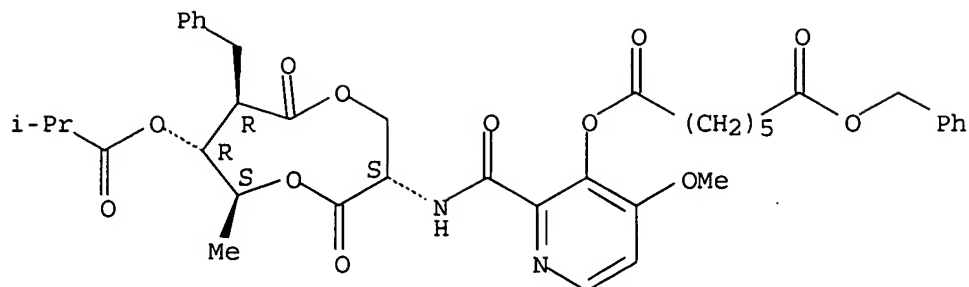


RN 234113-05-4 CAPLUS

CN Heptanedioic acid, 4-methoxy-2-[[[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-1-oxopropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]amino]carbonyl]-3-pyridinyl phenylmethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

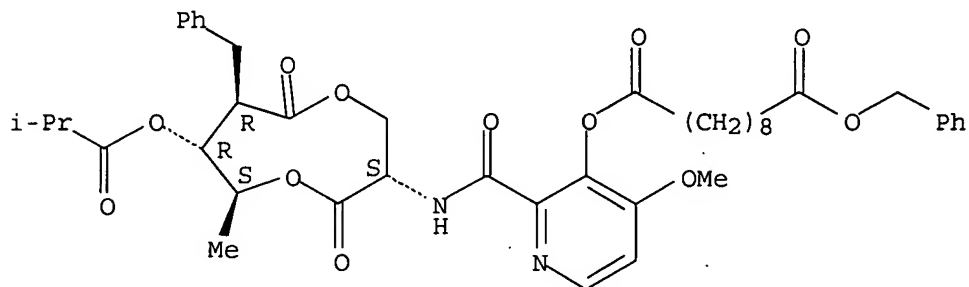
10/647,172



RN 234113-06-5 CAPLUS

CN Decanedioic acid, 4-methoxy-2-[[[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-1-oxopropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]amino]carbonyl]-3-pyridinyl phenylmethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.



IT 234112-91-5P 234112-93-7P 234112-94-8P
234112-95-9P 234112-96-0P 234112-97-1P
234112-98-2P 234112-99-3P 234113-00-9P
234113-01-0P 234113-02-1P 234113-03-2P
234113-04-3P 234113-07-6P 234113-08-7P
234113-09-8P 234113-10-1P 234113-11-2P
234113-12-3P 234113-30-5P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); IMF (Industrial manufacture); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

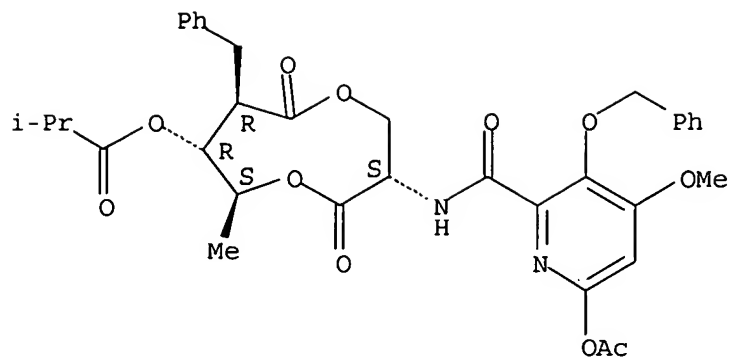
(preparation of UK-2A derivs. as antifungals)

RN 234112-91-5 CAPLUS

CN Propanoic acid, 2-methyl-, (3S,6S,7R,8R)-3-[[[6-(acetyloxy)-4-methoxy-3-(phenylmethoxy)-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

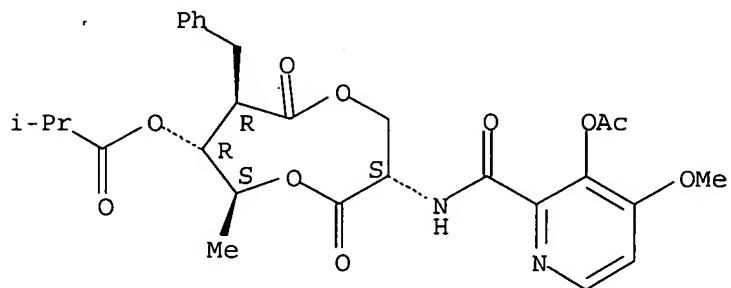
10/647,172



RN 234112-93-7 CAPLUS

CN Propanoic acid, 2-methyl-, (3S,6S,7R,8R)-3-[[[3-(acetyloxy)-4-methoxy-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

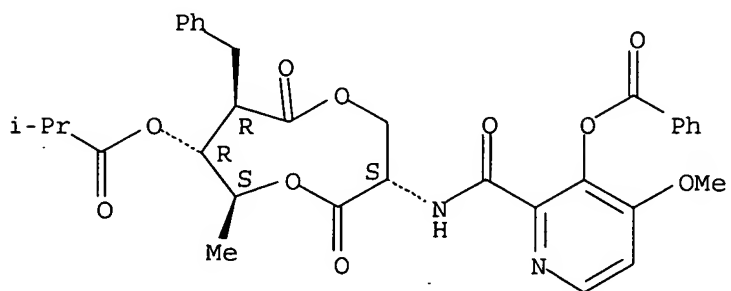
Absolute stereochemistry.



RN 234112-94-8 CAPLUS

CN Propanoic acid, 2-methyl-, (3S,6S,7R,8R)-3-[[[3-(benzoyloxy)-4-methoxy-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

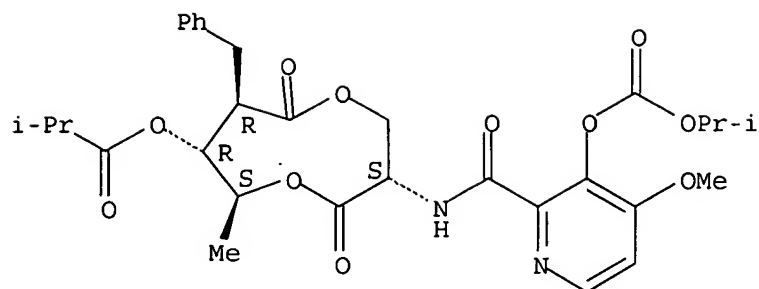


RN 234112-95-9 CAPLUS

CN Propanoic acid, 2-methyl-, (3S,6S,7R,8R)-3-[[[4-methoxy-3-[[[(1-methylethoxy)carbonyl]oxy]-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

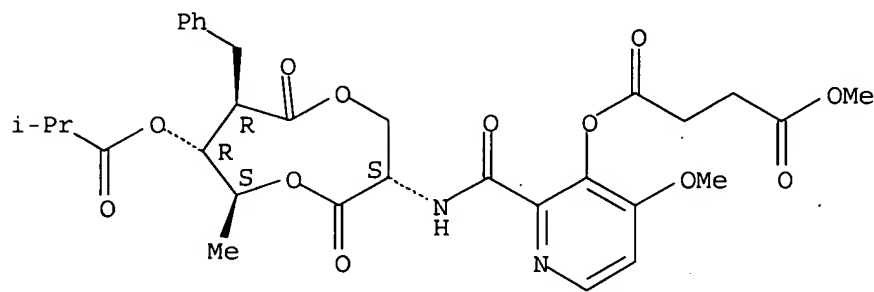
10/647,172



RN 234112-96-0 CAPLUS

CN Butanedioic acid, 4-methoxy-2-[[[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-1-oxopropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]amino]carbonyl]-3-pyridinyl methyl ester (9CI) (CA INDEX NAME)

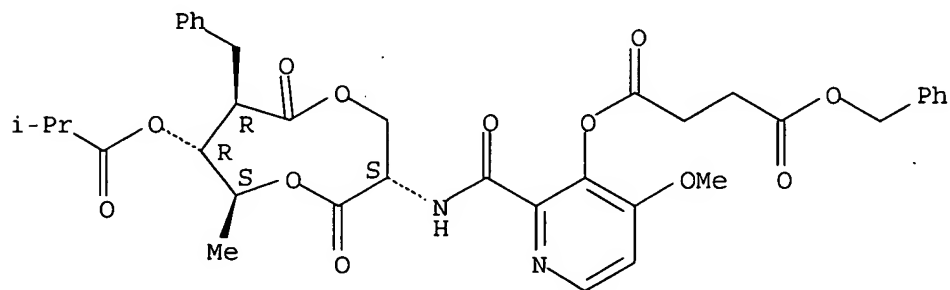
Absolute stereochemistry.



RN 234112-97-1 CAPLUS

CN Butanedioic acid, 4-methoxy-2-[[[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-1-oxopropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]amino]carbonyl]-3-pyridinyl phenylmethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

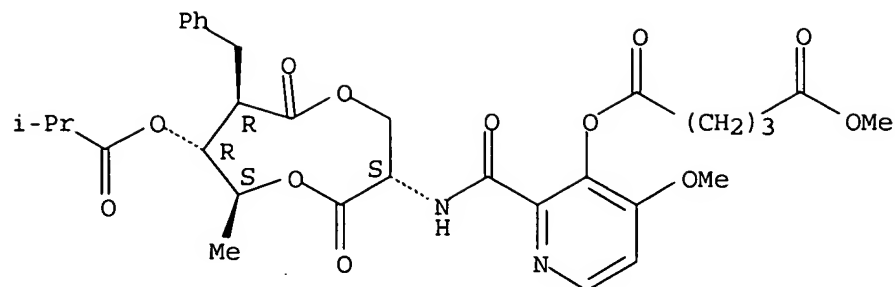


RN 234112-98-2 CAPLUS

CN Pentanedioic acid, 4-methoxy-2-[[[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-1-oxopropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]amino]carbonyl]-3-pyridinyl methyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

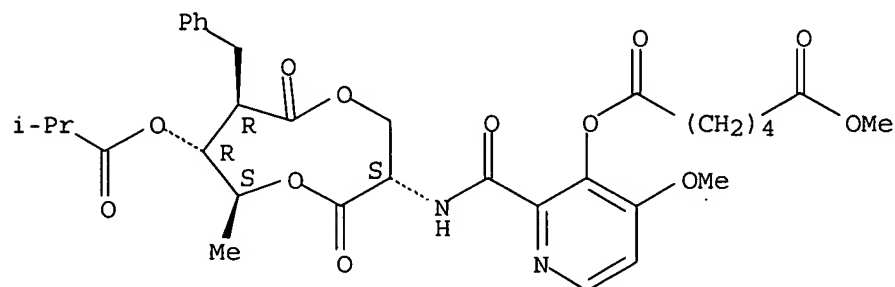
10/647,172



RN 234112-99-3 CAPLUS

CN Hexanedioic acid, 4-methoxy-2-[[[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-1-oxopropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]amino]carbonyl]-3-pyridinyl methyl ester (9CI) (CA INDEX NAME)

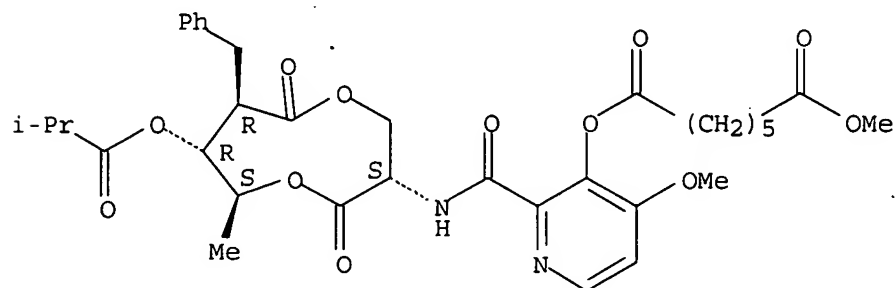
Absolute stereochemistry.



RN 234113-00-9 CAPLUS

CN Heptanedioic acid, 4-methoxy-2-[[[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-1-oxopropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]amino]carbonyl]-3-pyridinyl methyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

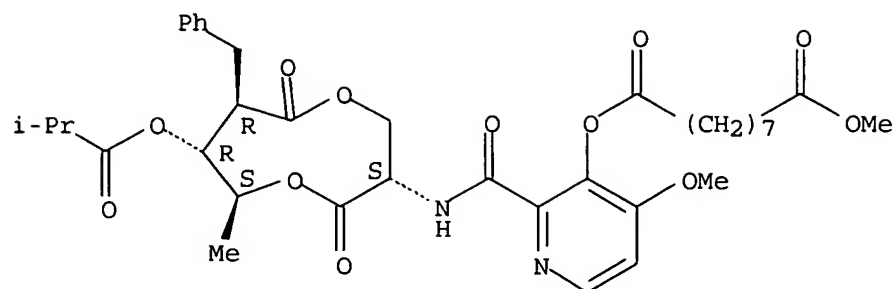


RN 234113-01-0 CAPLUS

CN Nonanedioic acid, 4-methoxy-2-[[[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-1-oxopropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]amino]carbonyl]-3-pyridinyl methyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

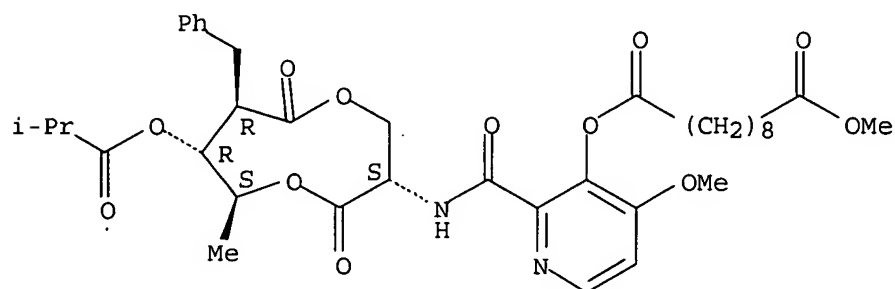
10/647,172



RN 234113-02-1 CAPLUS

CN Decanedioic acid, 4-methoxy-2-[[[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-1-oxopropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]amino]carbonyl]-3-pyridinyl methyl ester (9CI) (CA INDEX NAME)

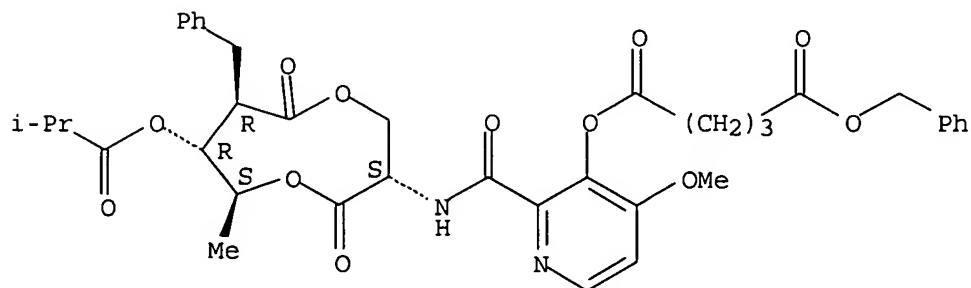
Absolute stereochemistry.



RN 234113-03-2 CAPLUS

CN Pentanedioic acid, 4-methoxy-2-[[[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-1-oxopropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]amino]carbonyl]-3-pyridinyl phenylmethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

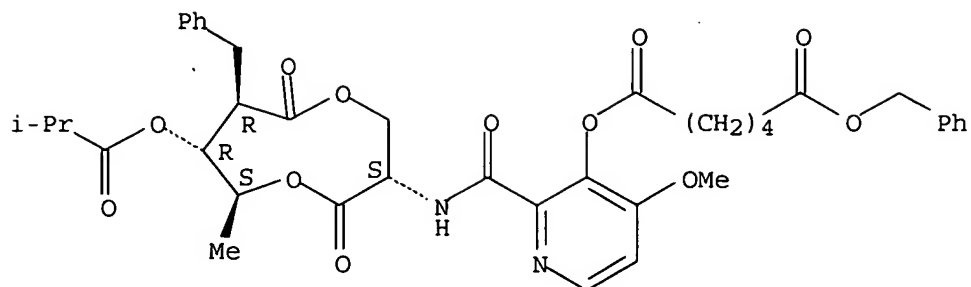


RN 234113-04-3 CAPLUS

CN Hexanedioic acid, 4-methoxy-2-[[[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-1-oxopropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]amino]carbonyl]-3-pyridinyl phenylmethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

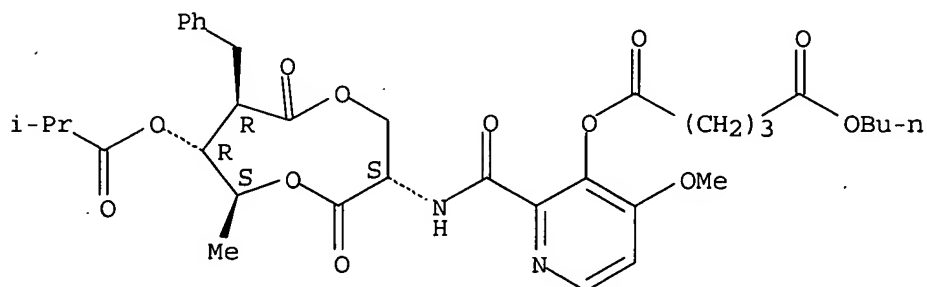
10/647,172



RN 234113-07-6 CAPLUS

CN Pentanedioic acid, butyl 4-methoxy-2-[[[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-1-oxopropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]amino]carbonyl]-3-pyridinyl ester (9CI) (CA INDEX NAME)

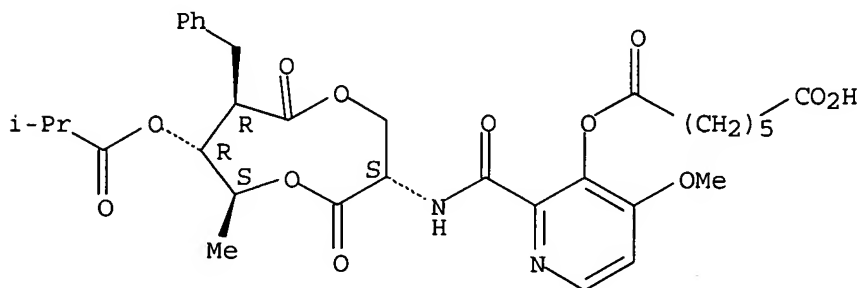
Absolute stereochemistry.



RN 234113-08-7 CAPLUS

CN Heptanedioic acid, mono[4-methoxy-2-[[[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-1-oxopropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]amino]carbonyl]-3-pyridinyl] ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

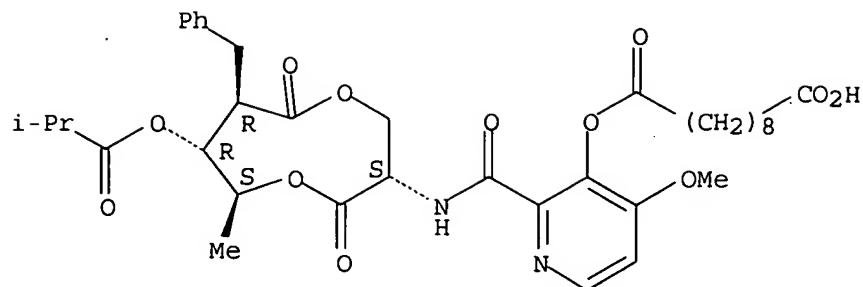


RN 234113-09-8 CAPLUS

CN Decanedioic acid, mono[4-methoxy-2-[[[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-1-oxopropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]amino]carbonyl]-3-pyridinyl] ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

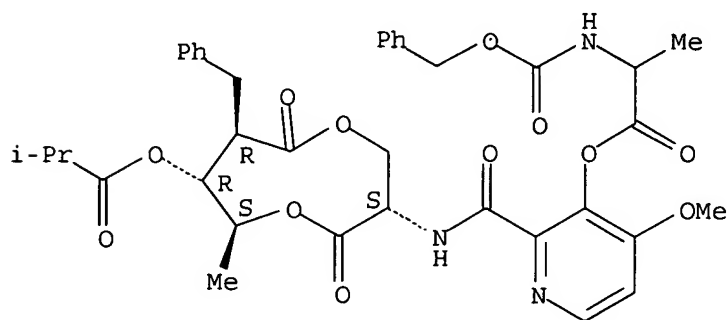
10/647,172



RN 234113-10-1 CAPLUS

CN Alanine, N-[(phenylmethoxy)carbonyl]-, 4-methoxy-2-[[[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-1-oxopropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]amino]carbonyl]-3-pyridinyl ester (9CI) (CA INDEX NAME)

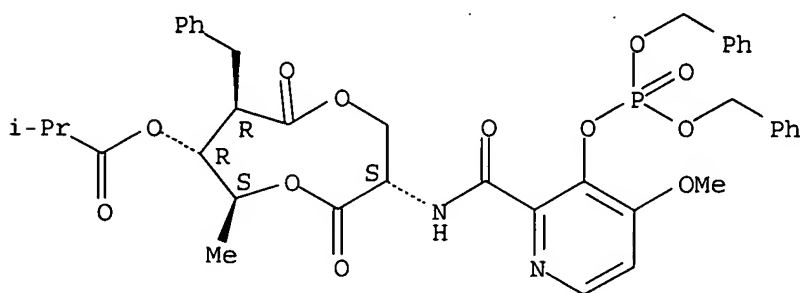
Absolute stereochemistry.



RN 234113-11-2 CAPLUS

CN Propanoic acid, 2-methyl-, (3S,6S,7R,8R)-3-[[[3-[[bis(phenylmethoxy)phosphinyl]oxy]-4-methoxy-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 234113-12-3 CAPLUS

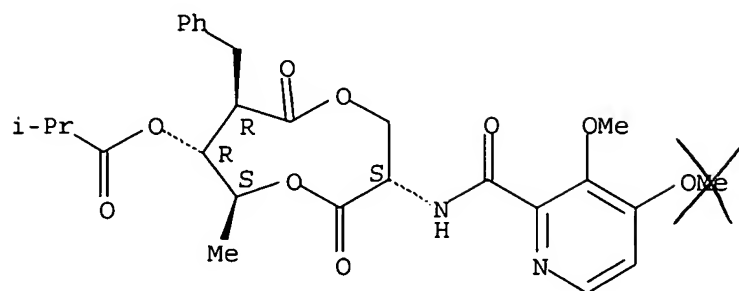
CN Propanoic acid, 2-methyl-, (3S,6S,7R,8R)-3-[[[3-[(diethoxyphosphinyl)oxy]-4-methoxy-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

10/647,172

[3S-(3R*,6R*,7S*,8S*)] - (9CI) (CA INDEX NAME)

Absolute stereochemistry.



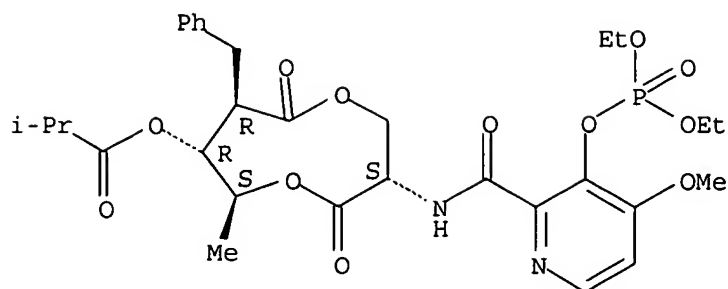
REFERENCE COUNT:

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THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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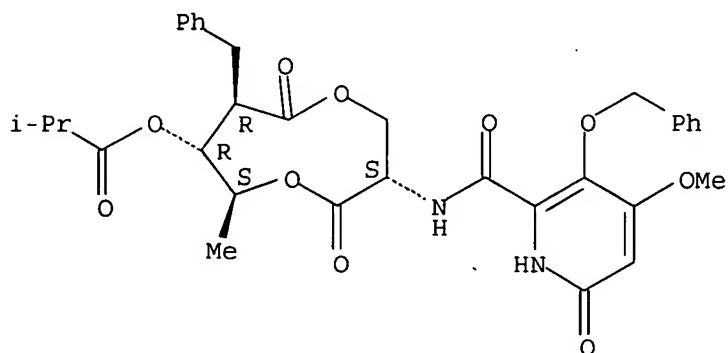
10/647,172



RN 234113-30-5 CAPLUS

CN Propanoic acid, 2-methyl-, (3S,6S,7R,8R)-3-[[[1,6-dihydro-4-methoxy-6-oxo-3-(phenylmethoxy)-2-pyridinyl]carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L19 ANSWER 7 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 1997:16443 CAPLUS

DOCUMENT NUMBER: 126:144017

TITLE: UK-2A, B, C and D, novel antifungal antibiotics from Streptomyces sp. 517-02. II. Structural elucidation

AUTHOR(S): Hanafi, Muhammad; Shibata, Kozo; Ueki, Masashi; Taniguchi, Makoto

CORPORATE SOURCE: Fac. Sci., Osaka City Univ., Osaka, 558, Japan

SOURCE: Journal of Antibiotics (1996), 49(12), 1226-1231

CODEN: JANTAJ; ISSN: 0021-8820

PUBLISHER: Japan Antibiotics Research Association

DOCUMENT TYPE: Journal

LANGUAGE: English

AB UK-2A, UK-2B, UK-2C and UK-2D, novel antibiotics produced by Streptomyces sp. 517-02, exhibit strong antifungal activity. The structures were elucidated based on spectral and chemical evidence that these compds. are the derivs. of the nine-membered dilactone formed from serine and 4-hydroxypentanoic acid moiety.

IT 186528-19-8P, O-Methyl UK 2A

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (structural elucidation of UK-2A, UK-2B, UK-2C and UK-2D, novel antifungal antibiotics from Streptomyces sp. 517-02)

RN 186528-19-8 CAPLUS

CN Propanoic acid, 2-methyl-, 3-[[[(3,4-dimethoxy-2-pyridinyl)carbonyl]amino]-6-methyl-4,9-dioxo-8-(phenylmethyl)-1,5-dioxonan-7-yl ester,